Physiological levels of thrombospondin-1 decrease NO-dependent vasodilation in coronary microvessels from aged rats

Chris Nevitt,1,2 Grant McKenzie,1 Katelyn Christian,1 Jeff Austin,1 Sarah Hencke,1 James Hoying,1,3 and Amanda LeBlanc1,3
1Cardiovascular Innovation Institute, University of Louisville, Louisville, Kentucky; 2Department of Biochemistry and Molecular Genetics, University of Louisville, Louisville, Kentucky; and 3Department of Physiology, University of Louisville, Louisville, Kentucky

Submitted 26 January 2016; accepted in final form 21 April 2016

Nevitt C, McKenzie G, Christian K, Austin J, Hencke S, Hoying J, LeBlanc A. Physiological levels of thrombospondin-1 decrease NO-dependent vasodilation in coronary microvessels from aged rats. Am J Physiol Heart Circ Physiol 310: H1842–H1850, 2016. First published May 3, 2016; doi:10.1152/ajpheart.00086.2016.—Aging and cardiovascular disease are associated with the loss of nitric oxide (NO) signaling and increases in oxidative stress, have been linked with aging (12, 21, 24, 25, 42). NO is the primary signaling molecule, and in particular, the arterioles whose vasoactivity is largely responsible for increasing myocardial perfusion to meet demand (15, 29). In addition, loss of NO signaling leads to a substrate switch of the proatherosclerotic molecule, H2O2 (6, 32, 33). Several studies have shown that a reduction in NO bioavailability contributes to the development of endothelial dysfunction (5, 39). Not surprisingly, loss of NO signaling and increases in oxidative stress are established markers of cardiovascular disease (9, 31, 33, 43, 45). Notably, endothelial dysfunction and loss of NO-mediated dilation have been shown to be effective predictors of future adverse cardiovascular events (2, 27).

The matricellular protein thrombospondin-1 (Thbs-1) has been shown to reduce NO-dependent vasodilation, with impact demonstrated on systemic blood pressure and in multiple vascular settings, including the aorta, kidney, and lung (3, 4, 10, 36). This effect occurs through inhibition of NO signaling at multiple levels, including blocking NO production through endothelial NO synthase (eNOS) and inhibiting the activity of soluble guanylyl cyclase (sGC), the major downstream effector of NO in vasodilation (3, 30, 46). Thbs-1 has additionally been shown to induce production of superoxide through NADPH oxidase and uncoupling of endothelial NO synthase (4, 10). Superoxide is known to rapidly react with NO to form peroxynitrite, reducing the availability of NO for vascular signaling and thereby pushing the balance of vasoactivity toward vasostriction (8). Binding to the cell surface receptor CD47 on both endothelial cells and vascular smooth muscle cells appears to facilitate the vascular actions of Thbs-1 (3, 10). In fact, the interaction of Thbs-1 and CD47 has been implicated in the progression of pulmonary arterial hypertension and renal ischemia-reperfusion injury (4, 36).

Because of the association with cardiovascular risk, it is important to understand the mechanisms by which NO signaling in the vasculature is impaired in aging, especially in women. This is because >50% of women that present with symptoms of coronary ischemia do not have large-vessel obstruction, but instead have coronary microvascular dysfunction (34). Our laboratory has previously demonstrated that NO-mediated dilation is decreased with advancing age in coronary arterioles of female rats (24). Thbs-1 is known to potently reduce NO signaling, but has not been studied in the context of coronary microvascular reactivity or advancing age in women. Understanding this process is likely to identify therapeutic targets for addressing coronary dysfunction in aging and the rationale for the use of female rodents in the present study. Additionally, peptides that mimic the CD47 binding domain of Thbs-1 have shown efficacy as anti-angiogenic therapies (17).
Humanized antibodies against CD47 have shown promise for cancer treatment in a preclinical setting and have progressed on to clinical trials (28). We hypothesize that Thbs-1 interaction with CD47 contributes to the loss of NO signaling with aging, and that use of antibodies designed to block the activation of CD47 will be effective for the treatment of age-related coronary microvascular dysfunction.

METHODS

Animals. All animal surgeries were performed in accordance with protocols approved by the University of Louisville Institutional Animal Care and Use Committee and the National Institutes of Health Guide for the Care and Use of Laboratory Animals (8th edition, 2011). All animals used were female Fischer 344 rats. Young rats were purchased from Harlan Laboratories and were used at 4 mo of age. Old rats were provided by the National Institute on Aging and were used at 24 mo of age. All animals were anesthetized with 5% isoflurane before being euthanized by removal of the heart.

Isolated vessel experiments. Coronary arterioles (100–150 μm diameter) branching from the left anterior descending artery (LAD) were isolated and cannulated between pipettes in a microvascular perfusion chamber (isolated vessel systems). Vessels were maintained at 37°C in a 3-ml physiological saline solution (PSS) bath containing Ca2+ Ringer solution, MOPS, NaH2PO4, glucose, pyruvic acid, EDTA, and bovine serum albumin. PSS was changed every 20 min to refresh substrates required for vasoactivity. Pressure across the vessel was stabilized at 45 mmHg. The luminal vessel diameter was measured with a video caliper system (Colorado Video Systems). Pressure and diameter measurements were recorded throughout the experiment with LabScribe3 software (iWorx). Vasodilation experiments were begun once vessels reached and maintained a spontaneous tone of 20% or greater, spontaneous tone = 100 × (maximum diameter - resting diameter)/maximum diameter. Vessels that failed to reach a spontaneous tone of 20% or greater were not used for experiments. Maximum dilation was measured after 30-min incubation in Ca2+-free PSS, followed by addition of sodium nitroprusside.

NO-dependent dilation was induced by addition of the NO donor diethylamine NONOate (DEA-NONO-ate, Cayman Chemical) to the vessel perfusion bath in 2-min increments to cover a concentration range from 1 × 10^-7 to 1 × 10^-4 M in half-log increments. Vessel diameter at the end of each 2-min incubation was recorded. In dilation experiments involving Thbs-1, vessels were incubated with 2.2 nM Thbs-1 (recombinant human, Athens Research and Technology) for 1 h before as well as during the concentration response of DEA-NONO-ate. Vessels were incubated with the SOD mimetic 4-hydroxy-2,2,6,6-tetramethylpiperidine 1-oxyl (Tempol, 0.1 mM, Sigma) and the enzy- cme polyethylene glycol-catalase (catalase, 500 U/ml, Sigma), along with 2 × 10^-4 M in half-log increments. Vessel diameter at the end of each 2-min incubation was recorded. In dilation experiments involving Thbs-1, vessels were incubated with 2.2 nM Thbs-1 (recombinant human, Athens Research and Technology) for 1 h before as well as during the concentration response of DEA-NONO-ate. Vessels were incubated with the SOD mimetic 4-hydroxy-2,2,6,6-tetramethylpiperidine 1-oxyl (Tempol, 0.1 mM, Sigma) and the enzyme polyethylene glycol-catalase (catalase, 500 U/ml, Sigma), along with CD47 transcript levels were determined by a Bradford Protein Quantification assay (Bio-Rad). Lysis buffer was used to bring all samples to a protein concentration of 1 μg/μl. Fifteen micrograms of each sample were prepared using 2 x Laemmlı buffer (Bio-Rad) containing 5% β-mercaptoethanol. Electrophoresis was performed using precast 4–20% gradient gels (Bio-Rad), followed by overnight transfer to polyvinylidene difluoride membranes. Membranes were blocked in 5% milk before overnight incubation with primary antibodies against Thbs-1 (mouse anti-Thbs-1, 1:200, Thermo Scientific), CD47 (goat anti-CD47, 1:1,000, Santa Cruz), and β-actin (mouse anti-β-actin, 1:2,000, Santa Cruz). Incubation with the fluorescently labeled secondary antibodies goat anti-mouse 488 and rabbit anti-goat 594 (Jackson ImmunoResearch) were carried out for 1 h at 1:1,000 dilutions. Imaging was carried out on a Typhoon FLA 7000 biomolecular imager (GE). ImageJ 1.47v software (National Institutes of Health) was used to perform densitometry, with β-actin serving as a loading control for each sample. Quantitative PCR. Thbs-1 and CD47 transcript levels were detected in LV tissue and coronary arterioles. RNA was isolated from LV tissue using RNA-bee (Amsbio) and phase-lock gel tubes (5 Prime). A RNeasy Mini Kit (Qiagen) was used according to the manufacturer’s protocols to isolate RNA from coronary arterioles. Following RNA isolation, cDNA was synthesized using the Affinity Script quantitative PCR (qPCR) cDNA synthesis kit (Agilent), according to manufacturer instructions. Quantitative real-time PCR was performed on the generated complimentary DNA to measure Thbs-1 and CD47 transcript levels, with readings being normalized to the reference gene dynactin. All reactions were run on a Rotor Gene 6000 real-time PCR machine (Corbett Life Science). The following rat primers (Integrated DNA Technologies) were used: Thbs-1 forward CCGTTTGATCAGATGGTGA, Thbs-1 reverse CGGCACCTGC- ATATGATACGT, CD47 forward CTCGGCCTGATGCCTGGCT, CD47 reverse CTGGTTGAAGCAGAAAGAC, dynactin forward CGAAGAGCTCAGATAGAGG, and dynactin reverse GAAGGT- CACTTTGCCCATGT. Fold changes were determined by the 2^(-ΔΔCT) method, in which ΔΔCT = ΔCT(target sample) - ΔCT(reference sample).

Immunofluorescence and confocal imaging. CD47 expression and localization within isolated coronary arterioles was analyzed by immunofluorescent labeling and confocal imaging. Coronary arterioles of 100- to 150-μm diameter were isolated from the LAD, fixed with 2% paraformaldehyde (Electron Microscopy Sciences), and permeabilized with 0.5% Triton X-100 (Sigma Aldrich). Blocking was carried in PBS containing 0.1% Triton X-100, 10% normal rabbit

AJP-Heart Circ Physiol • doi:10.1152/ajpheart.00086.2016 • www.ajpheart.org
CD47 blockade improves NO-dependent dilation in aging.

In a separate set of experiments, coronary arterioles from young and old female rats were cotreated with Thbs-1 and αCD47. Arterioles from young rats did not show differences in vasodilation with αCD47 treatment compared with Thbs-1 alone or no treatment (Fig. 1, C and E). However, αCD47 treatment improved NO-dependent vasodilation in the presence of Thbs-1 in coronary arterioles from old rats, with significant improvements at \( 1 \times 10^{-5} \) M (percent relaxation, Thbs-1 alone 16.28 ± 8.16% and Thbs-1 + αCD47 71.55 ± 8.33%), 3 \( \times 10^{-5} \) M (59.06 ± 3.58 and 77.34 ± 7.24%), and 1 \( \times 10^{-4} \) M (65.48 ± 2.80 and 84.02 ± 6.01%, respectively) DEA-NONO-ate (Fig. 1, D and 1F). Additionally, arterioles from old rats treated with αCD47 showed significantly increased dilation to low concentrations of NO compared with untreated arterioles (3 \( \times 10^{-7} \) M, untreated 4.56 ± 9.16% and Thbs-1 + αCD47 19.10 ± 7.16%; 1 \( \times 10^{-6} \) M, 4.06 ± 5.98 and 23.13 ± 6.57%, respectively) (compare Fig. 1, B and D). Notably, cotreatment of old arterioles with αCD47 and Thbs-1 improved vasodilation across all concentrations of DEA-NONO-ate to levels equal or greater than those seen in young arterioles under basal conditions (compare Fig. 1, A and D).

ROS scavengers minimally restore NO-dependent vasodilation in aging. Since Thbs-1 has been shown to increase superoxide levels, the SOD mimetic Tempol and the enzyme catalase were employed to convert superoxide to oxygen and water (4, 10). Arterioles from young rats cotreated with Thbs-1, Tempol, and catalase showed no difference in NO-dependent relaxation compared with Thbs-1 only and untreated vessels (Fig. 1, C and E). Cotreatment of coronary arterioles from old rats with Thbs-1, Tempol, and catalase improved NO-dependent vasodilation at only the 1 \( \times 10^{-4} \) M concentration of

---

**Table 1. Age group characteristics**

<table>
<thead>
<tr>
<th>Age Group</th>
<th>Young Female</th>
<th>Old Female</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mo</td>
<td>4</td>
<td>24</td>
</tr>
<tr>
<td>Total n</td>
<td>28</td>
<td>35</td>
</tr>
<tr>
<td>Body weight, g</td>
<td>194.9 ± 2.5</td>
<td>291.4 ± 6.8*</td>
</tr>
<tr>
<td>Heart weight, mg</td>
<td>510.7 ± 10.2</td>
<td>727.9 ± 16.8*</td>
</tr>
<tr>
<td>Maximum vessel diameter, μm</td>
<td>137.0 ± 5.0</td>
<td>143.2 ± 4.6</td>
</tr>
</tbody>
</table>

**RESULTS**

Thbs-1 reduces NO-dependent dilation in coronary arterioles of aged rats. Under basal conditions (untreated at an intravascular pressure of 45 mmHg), coronary arterioles from young (4 mo) and old (24 mo) female rats showed similar levels of maximal diameter and spontaneous tone (Table 1) and relaxation to DEA-NONO-ate (Fig. 1, A and B). The addition of exogenous Thbs-1, at physiologically relevant levels (10), into the vessel bath did not inhibit NO-dependent dilation in arterioles from young rats (Fig. 1A). In contrast, the presence of exogenous Thbs-1 led to significant impairment of NO-dependent dilation in arterioles from old rats at 3 \( \times 10^{-6} \) M (percent relaxation, untreated 47.32 ± 11.85% and Thbs-1 16.34 ± 10.33%), 1 \( \times 10^{-5} \) M (54.22 ± 11.20 and 16.28 ± 8.16%), 3 \( \times 10^{-5} \) M (72.25 ± 8.34 and 59.06 ± 3.58%), and 1 \( \times 10^{-4} \) M (84.96 ± 2.39 and 65.48 ± 2.80%, respectively) DEA-NONO-ate (Fig. 1B).

---

**Statistics.** Vessel relaxation and superoxide experiments were analyzed using fixed effects, mixed-model ANOVA. Bonferroni post hoc analysis was used to make subsequent pairwise comparisons. Experiments directly evaluating age-related differences without multiple treatments were analyzed by a Student’s t-test. In all cases, the threshold for significance was set to \( P \leq 0.05 \). All statistical analysis was performed using either SPSS (IBM) or SigmaPlot v11.2 software packages (Systat Software).

**CD47 blockade improves NO-dependent dilation in aging.** In a separate set of experiments, coronary arterioles from young and old female rats were cotreated with Thbs-1 and αCD47. Arterioles from young rats did not show differences in vasodilation with αCD47 treatment compared with Thbs-1 alone or no treatment (Fig. 1, C and E). However, αCD47 treatment improved NO-dependent vasodilation in the presence of Thbs-1 in coronary arterioles from old rats, with significant improvements at 1 \( \times 10^{-5} \) M (percent relaxation, Thbs-1 alone 16.28 ± 8.16% and Thbs-1 + αCD47 71.55 ± 8.33%), 3 \( \times 10^{-5} \) M (59.06 ± 3.58 and 77.34 ± 7.24%), and 1 \( \times 10^{-4} \) M (65.48 ± 2.80 and 84.02 ± 6.01%, respectively) DEA-NONO-ate (Fig. 1, D and 1F). Additionally, arterioles from old rats treated with αCD47 showed significantly increased dilation to low concentrations of NO compared with untreated arterioles (3 \( \times 10^{-7} \) M, untreated 4.56 ± 9.16% and Thbs-1 + αCD47 19.10 ± 7.16%; 1 \( \times 10^{-6} \) M, 4.06 ± 5.98 and 23.13 ± 6.57%, respectively) (compare Fig. 1, B and D). Notably, cotreatment of old arterioles with αCD47 and Thbs-1 improved vasodilation across all concentrations of DEA-NONO-ate to levels equal or greater than those seen in young arterioles under basal conditions (compare Fig. 1, A and D).

**ROS scavengers minimally restore NO-dependent vasodilation in aging.** Since Thbs-1 has been shown to increase superoxide levels, the SOD mimetic Tempol and the enzyme catalase were employed to convert superoxide to oxygen and water (4, 10). Arterioles from young rats cotreated with Thbs-1, Tempol, and catalase showed no difference in NO-dependent relaxation compared with Thbs-1 only and untreated vessels (Fig. 1, C and E). Cotreatment of coronary arterioles from old rats with Thbs-1, Tempol, and catalase improved NO-dependent vasodilation at only the 1 \( \times 10^{-4} \) M concentration of

---

**Values are means ± SE; n, no. of rats. Animal and isolated coronary arteriole characteristics are presented for young and old age female groups. Old female rats exhibited increased body weight and heart weight compared with young females. Initial steady-state diameter values were measured for each treatment before addition of DEA-NONO-ate, but no age-related or treatment-dependent differences were noted.**
Superoxide levels, as measured by DHE fluorescence intensity change, were similar for arterioles from both young and old rats, with increases observed after addition of Thbs-1 (Fig. 2B). Notably, Thbs-1 induced significantly greater superoxide production in arterioles from old rats compared with young (4 mo) rats. Addition of Tempol and catalase improved vasodilation compared with Thbs-1 alone at $1 \times 10^{-4}$ M DEA-NONO-ate.

CD47 blockade prevents Thbs-1-induced production of superoxide. Superoxide levels, as measured by DHE fluorescence intensity, were similar for arterioles from both young and old rats, with increases observed after addition of Thbs-1 (Fig. 2). Notably, Thbs-1 induced significantly greater superoxide production in arterioles from old rats compared with young [Fig. 2B; fluorescence intensity change, young $12.14 \pm 1.60$ and old $20.68 \pm 2.72$ arbitrary units (AU)]. As expected, incubation with a combination of Thbs-1, Tempol, and catalase prevented increases in superoxide levels, resulting in a significant difference compared with Thbs-1 alone for both young (Thbs-1 $12.14 \pm 1.60$ and Thbs-1 + Tempol and catalase $3.85 \pm 1.70$ AU) and old (20.68 $\pm 2.72$ and 4.21 $\pm 1.72$ AU), respectively) rats (Fig. 2). Suppression of superoxide production was also achieved when $\alpha$CD47 was provided along with Thbs-1 in both young (Thbs-1 $12.14 \pm 1.60$ and Thbs-1 + $\alpha$CD47 $2.02 \pm 1.99$ AU) and old (20.68 $\pm 2.72$ and 3.54 $\pm 2.18$ AU, respectively) rats (Fig. 2).

Intravenous administration of $\alpha$CD47 improves CFR in aged rats. Microsphere blood tracers were used to measure BF under baseline (resting) and dobutamine-stimulated (working) conditions in different regions of the heart. Twenty-four-
month-old rats receiving intravenous administration of αCD47 showed increased CFR (%increase in BF with dobutamine over baseline) in heart sections supplied by the LAD compared with rats receiving control IgG. Significant improvements in CFR were seen in the LV 1 (%increase, IgG 9.7 ± 9.3 and αCD47 84.0 ± 23.2%) and LV 2 (IgG 7.3 ± 9.9 and αCD47 67.5 ± 15.7%, respectively) heart sections (Fig. 5A). Absolute BF (ml·min⁻¹·g⁻¹) did not significantly differ between baseline and dobutamine conditions in any sections of the heart for control IgG-treated rats. In contrast, significant differences between baseline and dobutamine BF for αCD47-treated rats were seen in the apex (BF, baseline 1.45 ± 0.28 and dobutamine 2.84 ± 0.45 ml·min⁻¹·g⁻¹), LV 1 (baseline 1.85 ± 0.30 and dobutamine 3.24 ± 0.52 ml·min⁻¹·g⁻¹), and LV 2 (baseline 1.94 ± 0.29 and dobutamine 3.08 ± 0.31 ml·min⁻¹·g⁻¹) heart sections. Additionally, a significant difference was approached in the LV 3 (P = 0.06) heart section of αCD47-treated rats. Notably, absolute BF measurements did not significantly differ between control IgG and αCD47-treated rats, except in the apex section at baseline (IgG 2.60 ± 0.26 and αCD47 1.45 ± 0.28 ml·min⁻¹·g⁻¹) (Fig. 5B).

**DISCUSSION**

The mechanisms by which aging contributes to vascular dysfunction and loss of NO signaling are incompletely understood. Our findings show that there is an age-dependent dysfunction in coronary arterioles related to Thbs1-CD47 activity, leading to decreased vasodilation to NO and increased superoxide production within the coronary microvessel wall. Exogenous Thbs1 significantly decreased NO-dependent vasodilation in coronary arterioles from rats of advanced age (Fig. 1, B and F), while coronary arterioles from young rats treated with Thbs1 did not show altered NO responsiveness (Fig. 1, A and E). These data suggest that aging increases the response of the coronary microvessel wall to Thbs1, resulting in suppression of NO-dependent vasodilation (Fig. 6). This raises the possibility that Thbs1 is a contributor to the underlying coronary vascular pathology associated with aging.

Given the growing body of literature showing that Thbs1 suppresses NO signaling in general (3, 30, 46), it is somewhat surprising that coronary arterioles from young rats did not also show impaired NO-dependent vasodilation. A possibility is that healthy coronary arterioles are protected from the action of Thbs1 through mechanisms such as antioxidant defense, and that this protection is lost in aging. In fact, impaired antioxidant defenses are known contributors to the development of vascular dysfunction in aging (1, 9, 11, 12, 14). To investigate this possibility, the effect of aging on superoxide production downstream of Thbs1 was evaluated. Our results show that advanced age predisposes coronary arterioles to higher levels of superoxide production when challenged with Thbs1 (Fig. 2). We originally hypothesized that the increased presence of superoxide was reducing NO bioavailability and driving age-dependent inhibition of vasodilation by Thbs1. However, treating vessels with the antioxidants Tempol and catalase to reduce superoxide levels only minimally rescued vasodilation in arterioles from aged rats (Fig. 1, D and F). These results argue that superoxide generation is not the primary mechanism by which Thbs1 suppresses NO-dependent vasodilation in aging coronary arterioles. Thbs1 has previously been shown to directly inhibit sGC, and this represents a non-superoxide-driven pathway by which Thbs1 could be exerting an age-dependent effect on vasodilation (30). Interestingly, decreased expression and activity of sGC have previously been noted in advanced age (22, 37, 41).
Both suppression of NO-dependent vasodilation and production of superoxide in the presence of Thbs-1 were attenuated by treatment with a CD47 blocking antibody in coronary arterioles of old rats (Figs. 2 and 1, D and F). This adds to other reports that highlight CD47 as the necessary cell surface receptor for Thbs-1 regulation of NO signaling (3, 10, 18, 30). The cell surface receptor CD36 has also been described as a binding partner for Thbs-1, mediating many of the same cellular responses as CD47, particularly in the context of angiogenesis and platelet aggregation (13, 19, 20). However, inhibition of NO signaling by physiological concentrations of Thbs-1 is attenuated in CD47-null, but not CD36-null, models (18). This agrees with our results, which show that, in coronary arterioles, the effect of Thbs-1 on superoxide production and vasodilation can be counteracted by blocking CD47. As such, our work supports the targeting of CD47 for treating age-related vascular dysfunction.

The positive effect of αCD47 treatment on vasodilation seen in isolated vessel experiments was also observed in vivo. Aged rats treated with αCD47 showed improved CFR in sections of the heart supplied by the LAD and responsible for generating much of the contractile force of the LV (Fig. 5). Decreased CFR, associated with aging and cardiovascular disease, limits the ability of the heart to respond to increased demand and is associated with cardiovascular disease (16, 38). Blocking of CD47 appears to be a promising therapeutic strategy for increasing CFR and addressing age-related coronary microvascular dysfunction. Humanized CD47 blocking antibodies are currently in phase I clinical trials for cancer treatment (28). CD47 is frequently overexpressed by tumor cells, where it acts to suppress immune clearance through interaction with signal...
regulatory protein-α on myeloid cells (44). These trials will evaluate the safety and efficacy of CD47 blocking therapeutics, potentially bringing CD47 blockade as a therapy for vascular disease one step closer to reality.

The mechanism by which age contributes to increased coronary microvascular dysfunction downstream of Thbs-1 is not yet understood. In this study, Thbs-1 was provided at 2.2 nM to reflect reported circulating concentrations. Plasma levels of Thbs-1 for young and old female rats were both within error of this reported value, with concentrations found to be 1.72 and 0.95 nM, respectively (Fig. 4). Plasma, heart, and coronary arteriole levels of Thbs-1 also did not differ statistically based on age in our female rats (Figs. 3, A–C, and 4). Our study is the first to report Thbs-1 expression in young or old female rats. The lack of age-related changes in Thbs-1 expression contrasts with two previous reports that found Thbs-1 levels to increase with age in myocardial tissue and skin biopsies (7, 35). Both of these studies used male mice, whereas we have employed a female rat model, suggesting this difference may be related to sex or species. Decreased antioxidant defense is known to occur in aging and may play a role in increasing superoxide levels in coronary arterioles of old rats after treatment with Thbs-1 (9, 43). Increased superoxide levels alone do not account for the full age-dependent effect of Thbs-1, as scavenging of reactive oxygen species did not completely reverse

Fig. 5. CD47 blockade improves coronary blood flow reserve (CFR) in advanced age. A: CFR was calculated from measurements of blood flow at baseline and with dobutamine stimulation. Hearts from control IgG-treated (n = 4) and αCD47-treated (n = 5) 24-mo-old rats were sectioned to allow for analysis of regional perfusion changes. Color for each section corresponds to percent increase in blood flow with dobutamine stimulation over baseline. *Significant difference between αCD47 and control IgG sections. B: absolute blood flow (ml-min⁻¹·g⁻¹) under baseline and dobutamine conditions is shown for heart sections from control IgG- and αCD47-treated old rats. Values are means ± SE. *Significant difference between dobutamine stimulated and baseline blood flow. †Difference between control IgG and αCD47 treated rats.

Fig. 6. Pathways downstream of CD47 are implicated in the exacerbated response to Thbs-1 observed in coronary arterioles from aged rats. Coronary arterioles from old rats show increased superoxide production and decreased NO-dependent vasodilation in response to Thbs-1 compared with coronary arterioles from young rats. These effects are independent of Thbs-1 and CD47 concentration. Thbs-1 was added exogenously at equal levels to both age groups, and CD47 density on coronary arterioles was not found to be altered in aging. As such, more potent activation of CD47 and greater effect on downstream effectors are implicated in the exacerbated response to Thbs-1 seen in arterioles from aged rats. Downstream effects of CD47 include activation of NADPH oxidase, uncoupling of endothelial NO synthase (eNOS), and inhibition of soluble guanylyl cyclase (sGC).
Thbs1 effects on vasodilatation (Fig. 1D). Increased vascular density of CD47 in aging would contribute to increased Thbs-1 binding events and signal transduction, but qPCR and immunofluorescence of coronary artery arterioles did not show significant variation in CD47 receptor expression with age in females (Fig. 3, C and D). Strasbourg et al. (40) demonstrated that, in patients with metabolic syndrome, there was a significant decrease in CD47 in red blood cells from men compared with women, but no sex difference was found in cells from age-matched healthy donors. To date, there have been very few sex-specific comparisons of the Thbs-1/CD47 signaling pathway and expression as age progresses, which implies that this is an understudied research area. Our results suggest that, even without increases in concentration in either Thbs-1 or CD47 in our female rats, Thbs-1 induces increased vascular impairment in aging via enhanced CD47 activity.

We have shown that Thbs-1, at circulating levels typical of young rats, leads to suppression of NO-dependent vasodilatation and heightened superoxide production in advanced age. Regardless of the mechanism, age-dependent response to Thbs-1 may be a direct contributor to the development of a negative vascular phenotype in aging (Fig. 6), with consequences for cardiovascular health. Blockade of CD47 attenuates the effects of Thbs-1 and, therefore, represents a therapeutic target for age-related cardiovascular dysfunction that deserves further attention.

REFERENCES


42. Sverdlov AL, Ngo DT, Chan WP, Chirkov YY, Horowitz JD. Aging of the nitric oxide system: are we as old as our NO? J Am Heart Assoc. e000973, 2014.


